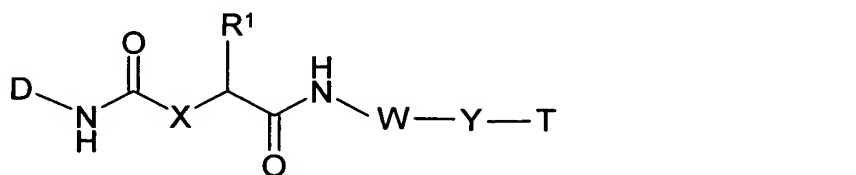


This listing of claims will replace all prior versions of claims in the application.

**Listing of Claims:** Please amend the claims as follows:

**We claim:**

**Claim 1. (Currently Amended)** ~~Compounds~~ A compound of the formula I



in which wherein

D denotes is an aromatic five-membered heterocyclic ring having 1 to 4 N, O and/or S atoms which is unsubstituted or mono- or polysubstituted by Hal, A, OR<sup>2</sup>, N(R<sup>2</sup>)<sub>2</sub>, NO<sub>2</sub>, CN, COOR<sup>2</sup> or CON(R<sup>2</sup>)<sub>2</sub>,

X denotes is NR<sup>3</sup> or O,

R<sup>1</sup> denotes is H, Ar, Het, cycloalkyl or A, which ~~may be~~ is optionally substituted by OR<sup>2</sup>, SR<sup>2</sup>, N(R<sup>2</sup>)<sub>2</sub>, Ar, Het, cycloalkyl, CN, COOR<sup>2</sup> or CON(R<sup>2</sup>)<sub>2</sub>,

R<sup>2</sup> denotes is H, A, -[C(R<sup>3</sup>)<sub>2</sub>]<sub>n</sub>-Ar, -[C(R<sup>3</sup>)<sub>2</sub>]<sub>n</sub>-Het, -[C(R<sup>3</sup>)<sub>2</sub>]<sub>n</sub>-cycloalkyl, -[C(R<sup>3</sup>)<sub>2</sub>]<sub>n</sub>-N(R<sup>3</sup>)<sub>2</sub> or -[C(R<sup>3</sup>)<sub>2</sub>]<sub>n</sub>-OR<sup>3</sup>,

R<sup>3</sup> denotes is H or A,

W denotes is -[C(R<sup>3</sup>)<sub>2</sub>]<sub>n</sub>-,

Y denotes is alkylene, cycloalkylene, Het-diyl or Ar-diyl,

T denotes is a mono- or bicyclic saturated, unsaturated or aromatic carbo- or heterocyclic ring having 0 to 4 N, O and/or S atoms, which ~~may be~~ is optionally unsubstituted or mono-, di- or trisubstituted by Hal, A, -[C(R<sup>3</sup>)<sub>2</sub>]<sub>n</sub>-Ar, -[C(R<sup>3</sup>)<sub>2</sub>]<sub>n</sub>-Het, -[C(R<sup>3</sup>)<sub>2</sub>]<sub>n</sub>-cycloalkyl, OR<sup>3</sup>, N(R<sup>3</sup>)<sub>2</sub>, NO<sub>2</sub>, CN, COOR<sup>2</sup>, CON(R<sup>2</sup>)<sub>2</sub>, NR<sup>2</sup>COA, NR<sup>2</sup>CON(R<sup>2</sup>)<sub>2</sub>, NR<sup>2</sup>SO<sub>2</sub>A, COR<sup>2</sup>, SO<sub>2</sub>NR<sup>2</sup> and/or S(O)<sub>m</sub>A and/or carbonyl oxygen,

or  $N(R^2)_2$

and, if Y = piperidine-1,4-diyl, also  $R^2$  or cycloalkyl,

A denotes is unbranched or branched alkyl having 1-10 C atoms, in which wherein one or two  $CH_2$  groups ~~may be~~ are optionally replaced by O or S atoms and/or by  $-CH=CH-$  groups and/or also wherein 1-7 H atoms ~~may be~~ are optionally replaced by F,

Ar denotes is phenyl, naphthyl or biphenyl, each of which is, independently of one another, unsubstituted or mono-, di- or trisubstituted by Hal, A,  $OR^3$ ,  $N(R^3)_2$ ,  $NO_2$ , CN,  $COOR^3$ ,  $CON(R^3)_2$ ,  $NR^3COA$ ,  $NR^3CON(R^3)_2$ ,  $NR^3SO_2A$ ,  $COR^3$ ,  $SO_2N(R^3)_2$ ,  $S(O)_mA$ ,  $-[C(R^3)_2]_n-COOR^{2'}$  or  $-O-[C(R^3)_2]_o-COOR^{2'}$ ,

$R^{2'}$  denotes is H, A,  $-[C(R^3)_2]_n-Ar'$ ,  $-[C(R^3)_2]_n-Het'$ ,  $-[C(R^3)_2]_n-cycloalkyl$ ,  $-[C(R^3)_2]_n-N(R^3)_2$  or  $-[C(R^3)_2]_n-OR^3$ ,

$R^{2''}$  denotes is H, A,  $-[C(R^3)_2]_n-Ar'$  or  $-[C(R^3)_2]_n-cycloalkyl$ ,  $-[C(R^3)_2]_n-N(R^3)_2$  or  $-[C(R^3)_2]_n-OR^3$ ,

Ar' denotes is phenyl or benzyl, each of which is, independently of one another, unsubstituted or mono- or disubstituted by Hal or A,

Het denotes is a mono- or bicyclic saturated, unsaturated or aromatic heterocyclic ring having 1 to 4 N, O and/or S atoms, which ~~may be~~ is unsubstituted or mono-, di- or trisubstituted by carbonyl oxygen,  $=S$ ,  $=N(R^3)_2$ , Hal, A,  $-[C(R^3)_2]_n-Ar$ ,  $-[C(R^3)_2]_n-Het^1$ ,  $-[C(R^3)_2]_n-cycloalkyl$ ,  $-[C(R^3)_2]_n-OR^{2'}$ ,  $-[C(R^3)_2]_n-N(R^2)_2$ ,  $NO_2$ , CN,  $-[C(R^3)_2]_n-COOR^{2'}$ ,  $-[C(R^3)_2]_n-CON(R^2)_2$ ,  $-[C(R^3)_2]_n-NR^{2'}COA$ ,  $NR^{2'}CON(R^2)_2$ ,  $-[C(R^3)_2]_n-NR^{2'}SO_2A$ ,  $COR^{2'}$ ,  $SO_2NR^{2'}$  and/or  $S(O)_mA$ ,

Het<sup>1</sup> denotes is a mono- or bicyclic saturated, unsaturated or aromatic heterocyclic ring having 1 to 2 N, O and/or S atoms, which ~~may be~~ is unsubstituted or mono- or disubstituted by carbonyl oxygen,  $=S$ ,  $=N(R^3)_2$ , Hal, A,  $OR^{2''}$ ,  $N(R^{2''})_2$ ,  $NO_2$ , CN,  $COOR^{2''}$ ,  $CON(R^{2''})_2$ ,  $NR^{2''}COA$ ,  $NR^{2''}CON(R^{2''})_2$ ,  $NR^{2''}SO_2A$ ,  $COR^{2''}$ ,  $SO_2NR^{2''}$  and/or  $S(O)_mA$ ,

Hal denotes is F, Cl, Br or I,

n denotes is 0, 1 or 2,

m denotes is 0, 1 or 2,

o denotes is 1, 2 or 3,  
and or a pharmaceutically usable derivatives acceptable salt, solvate[s]  
and or stereoisomer[s] thereof, including or a mixture[s] thereof in all  
ratios.

**Claim 2. (Currently Amended)** Compounds A compound according to Claim 1, in  
which wherein

D denotes is an aromatic five-membered heterocyclic ring having 1  
to 2 N, O and/or S atoms which is unsubstituted or mono- or  
disubstituted by Hal,  
and or a pharmaceutically usable derivatives acceptable salt, solvate[s] and or  
stereoisomer[s] thereof, including or a mixture[s] thereof in all ratios.

**Claim 3. (Currently Amended)** Compounds A compound according to Claim 1, in  
which wherein

D denotes is a thienyl ring which is mono- or disubstituted by Hal,  
and or a pharmaceutically usable derivatives acceptable salt, solvate[s] and or  
stereoisomer[s] thereof, including or a mixture[s] thereof in all ratios.

**Claim 4. (Currently Amended)** Compounds A compound according to Claim 1, in  
which wherein

R<sup>2</sup> denotes is H or alkyl having 1, 2, 3, 4, 5 or 6 C atoms,  
and or a pharmaceutically usable derivatives acceptable salt, solvate[s] and or  
stereoisomer[s] thereof, including or a mixture[s] thereof in all ratios.

**Claim 5. (Currently Amended)** Compounds A compound according to Claim 1, in  
which wherein

R<sup>1</sup> denotes is H or unsubstituted phenyl, thienyl or alkyl having 1-6  
C atoms,  
and or a pharmaceutically usable derivatives acceptable salt, solvate[s] and or  
stereoisomer[s] thereof, including or a mixture[s] thereof in all ratios.

**Claim 6. (Currently Amended)** ~~Compounds~~ A compound according to Claim 1, in which wherein

X denotes is NH or O,

and or a pharmaceutically ~~usable derivatives~~ acceptable salt, solvate[s] and or stereoisomer[s] thereof, including or a mixture[s] thereof in all ratios.

**Claim 7. (Currently Amended)** ~~Compounds~~ A compound according to Claim 1, in which wherein

W denotes is (CH<sub>2</sub>)<sub>n</sub>,

and or a pharmaceutically ~~usable derivatives~~ acceptable salt, solvate[s] and or stereoisomer[s] thereof, including or a mixture[s] thereof in all ratios.

**Claim 8. (Currently Amended)** ~~Compounds~~ A compound according to Claim 1, in which wherein

Y denotes is Ar-diyl or Het-diyl,

and or a pharmaceutically ~~usable derivatives~~ acceptable salt, solvate[s] and or stereoisomer[s] thereof, including or a mixture[s] thereof in all ratios.

**Claim 9. (Currently Amended)** ~~Compounds~~ A compound according to Claim 1, in which wherein

T denotes is a mono- or bicyclic saturated, unsaturated or aromatic heterocyclic ring having 1 to 2 N and/or O atoms, which ~~may be~~ is unsubstituted or mono- or disubstituted by carbonyl oxygen, or N(R<sup>2</sup>)<sub>2</sub>

and, if Y = piperidine-1,4-diyl, also R<sup>2</sup>,

and or a pharmaceutically ~~usable derivatives~~ acceptable salt, solvate[s] and or stereoisomer[s] thereof, including or a mixture[s] thereof in all ratios.

**Claim 10. (Currently Amended)** ~~Compounds~~ A compound according to Claim 1, in which wherein

T denotes is a mono- or bicyclic saturated or unsaturated heterocyclic ring having 1 to 2 N and/or O atoms which is mono- or

disubstituted by carbonyl oxygen (=O),

or  $N(R^2)_2$

and, if Y = piperidine-1,4-diyl, also  $R^2$ ,

and or a pharmaceutically ~~usable derivatives~~ acceptable salt, solvate[s] and or stereoisomer[s] thereof, including or a mixture[s] thereof in all ratios.

**Claim 11. (Currently Amended)**

Compounds A compound according to Claim 1,

in which wherein

T denotes is piperidin-1-yl, pyrrolidin-1-yl, 1*H*-pyridin-1-yl, morpholin-4-yl, piperazin-1-yl, 1,3-oxazolidin-3-yl, 2*H*-pyridazin-2-yl, pyrazin-1-yl, azepan-1-yl, 2-azabicyclo[2.2.2]octan-2-yl, each of which is mono- or disubstituted by carbonyl oxygen, or  $N(R^2)_2$

and, if Y = piperidine-1,4-diyl, also  $R^2$ ,

and or a pharmaceutically ~~usable derivatives~~ acceptable salt, solvate[s] and or stereoisomer[s] thereof, including or a mixture[s] thereof in all ratios.

**Claim 12. (Currently Amended)**

Compounds A compound according to Claim 1,

in which wherein

Ar denotes is phenyl which is unsubstituted or mono- or disubstituted by Hal, A, OA,  $SO_2A$ ,  $COOR^2$ ,  $SO_2NH_2$  or CN,

and or a pharmaceutically ~~usable derivatives~~ acceptable salt, solvate[s] and or stereoisomer[s] thereof, including or a mixture[s] thereof in all ratios.

**Claim 13. (Currently Amended)**

Compounds A compound according to Claim 1,

in which wherein

D denotes is an aromatic five-membered heterocyclic ring having 1 to 2 N, O and/or S atoms which is unsubstituted or mono- or disubstituted by Hal,

$R^1$  denotes is H or unsubstituted phenyl, thienyl or alkyl having 1-6 C atoms,

$R^2$  denotes is H or alkyl having 1, 2, 3, 4, 5 or 6 C atoms,

X denotes NH or O,  
W denotes is W (CH<sub>2</sub>)<sub>n</sub>,  
Y denotes is Ar-diyl, pyridinediyl or piperidinediyl,  
Ar denotes is phenyl which is unsubstituted or mono- or disubstituted by Hal, A, OA, SO<sub>2</sub>A, COOR<sup>2</sup>, SO<sub>2</sub>NH<sub>2</sub> or CN,  
T denotes is piperidin-1-yl, pyrrolidin-1-yl, 1*H*-pyridin-1-yl, morpholin-4-yl, piperazin-1-yl, 1,3-oxazolidin-3-yl, 2*H*-pyridazin-2-yl, pyrazin-1-yl, azepan-1-yl, 2-azabicyclo[2.2.2]octan-2-yl, each of which is mono- or disubstituted by carbonyl oxygen, or N(R<sup>2</sup>)<sub>2</sub>  
and, if Y = piperidine-1,4-diyl, also R<sup>2</sup>,

and or a pharmaceutically usable derivatives acceptable salt, solvate[s] and or stereoisomer[s] thereof, ~~including or a~~ mixture[s] thereof in all ratios.

**Claim 14. (Currently Amended)**      Compounds A compound according to Claim 1, in which wherein

D denotes is thienyl, thiazolyl or furyl, each of which is mono- or disubstituted by Hal,  
R<sup>1</sup> denotes is H or unsubstituted phenyl, thienyl or alkyl having 1-6 C atoms,  
R<sup>2</sup> denotes is H or alkyl having 1, 2, 3, 4, 5 or 6 C atoms,  
X denotes is NH or O,  
W denotes is W (CH<sub>2</sub>)<sub>n</sub>,  
Y denotes is Ar-diyl, pyridinediyl or piperidinediyl,  
Ar denotes is phenyl which is unsubstituted or mono- or disubstituted by Hal, A, OA, SO<sub>2</sub>A, COOR<sup>2</sup>, SO<sub>2</sub>NH<sub>2</sub> or CN,  
T denotes is piperidin-1-yl, pyrrolidin-1-yl, pyridinyl, morpholin-4-yl, piperazin-1-yl, 1,3-oxazolidin-3-yl, pyridazin-2-yl, pyrazinyl, azepan-1-yl, 2-azabicyclo[2.2.2]octan-2-yl, each of which is unsubstituted or mono- or disubstituted by carbonyl oxygen, or N(R<sup>2</sup>)<sub>2</sub>  
and, if Y = piperidine-1,4-diyl, also R<sup>2</sup>,

and or a pharmaceutically usable ~~derivatives~~ acceptable salt, solvate[s] and or stereoisomer[s] thereof, ~~including or a~~ mixture[s] thereof in all ratios.

**Claim 15. (Currently Amended)**      ~~Compounds~~ A compound according to Claim 1 selected from the group which is

(R)-2-[3-(5-chlorothiophen-2-yl)ureido]-N-[4-(3-oxomorpholin-4-yl)-phenyl]valeramide,

(R)-2-[3-(5-chlorothiophen-2-yl)ureido]-N-[4-(3-oxomorpholin-4-yl)-3-methylphenyl]valeramide,

2-[3-(5-chlorothiophen-2-yl)ureido]-N-[4-(3-oxomorpholin-4-yl)-phenyl]acetamide,

(R)-2-[3-(5-bromothiophen-2-yl)ureido]-N-[4-(3-oxomorpholin-4-yl)-phenyl]valeramide,

(R)-2-[3-(5-bromofuran-2-yl)ureido]-N-[4-(3-oxomorpholin-4-yl)-phenyl]valeramide,

(R)-2-[3-(5-chlorothiophen-2-yl)ureido]-N-[4-(3-oxomorpholin-4-yl)-phenyl]-2-phenylacetamide,

(R)-2-[3-(5-chlorothiophen-2-yl)ureido]-N-[4-(3-oxomorpholin-4-yl)-phenyl]-2-(thiophen-2-yl)acetamide,

(R)-2-[3-(5-chlorothiophen-2-yl)ureido]-N-[4-(2-oxopiperidin-1-yl)-phenyl]valeramide,

(R)-2-[3-(5-chlorothiophen-2-yl)ureido]-N-[4-(2-oxo-1*H*-pyrazin-1-yl)-phenyl]valeramide,

(R)-2-[3-(5-chlorothiophen-2-yl)ureido]-N-[2-oxo-3,4,5,6-tetrahydro-[1,2']bipyridinyl-5'-yl]valeramide,

(S)-2-[3-(5-chlorothiophen-2-yl)ureido]-N-[4-(3-oxomorpholin-4-yl)-phenyl]-2-phenylacetamide,

(R)-2-[3-(5-chlorothiophen-2-yl)ureido]-N-[4-(3-oxomorpholin-4-yl)-phenylmethyl]valeramide,

(R)-2-[3-(5-chlorothiazol-2-yl)ureido]-N-[4-(3-oxomorpholin-4-yl)-phenyl]valeramide,

(R)-2-[N-(5-chlorothiophen-2-yl)carbamoyloxy]-N-[[4-(3-oxo-morpholin-4-yl)phenyl]valeramide,

(R)-2-[N-(5-chlorothiophen-2-yl)carbamoyloxy]-N-[C-(3,4,5,6-tetrahydro-2H-[1,4']bipyridinyl-4-yl)methyl]valeramide,

(R)-2-[N-(5-chlorothiophen-2-yl)carbamoyloxy]-N-[1-isopropyl-piperidin-4-ylmethyl]-2-phenylacetamide,

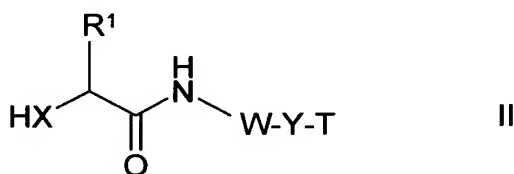
(R)-2-[N-(5-chlorothiophen-2-yl)carbamoyloxy]-N-[[4-(morpholin-4-yl)phenyl]valeramide; or

(R)-2-[N-(5-chlorothiophen-2-yl)carbamoyloxy]-N-(4-dimethylamino-phenyl)-2-phenylacetamide,

and or a pharmaceutically usable derivatives acceptable salt, solvate[s] and or stereoisomer[s] thereof, including or a mixture[s] thereof in all ratios.

**Claim 16. (Currently Amended)** ~~Process A~~ process for the preparation of compounds a compound of the formula I according to Claim 1 and pharmaceutically usable derivatives, solvates and stereoisomers thereof, characterised in that comprising

a) reacting a compound of the formula II



~~in which~~ wherein

R<sup>1</sup>, W, X, Y and T have the meaning indicated in Claim 1,

~~is reacted~~ with a compound of the formula III



~~in which~~ wherein

D has the meaning indicated in Claim 1,

or

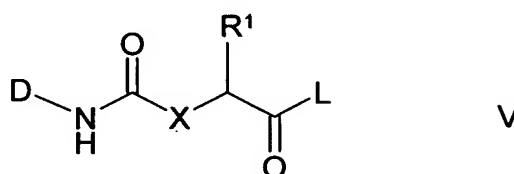


b) reacting a compound of the formula IV



~~in which~~ wherein W, Y and T have the meaning indicated in Claim 1,

~~is reacted~~ with a compound of the formula V



~~in which~~ wherein

L denotes Cl, Br, I or a free or reactively functionally modified OH group, and

R<sup>1</sup>, X and D have the meanings indicated in Claim 1,  
and/or

optionally converting a base or acid of the formula I ~~is converted~~ into one of its salts.

**Claim 17. (Currently Amended)**      Compounds of the formula I according to Claim 1 as inhibitors of A method of inhibiting coagulation factor Xa comprising contacting said coagulation factor Xa with a compound according to claim 1.

**Claim 18. (Currently Amended)**      Compounds of the formula I according to Claim 1 as inhibitors of A method of inhibiting coagulation factor VIIa comprising contacting said coagulation factor VIIa with a compound according to claim 1.

**Claim 19. (Currently Amended)**      Medicaments A pharmaceutical composition comprising at least one compound of the formula I according to Claim 1 ~~and/or pharmaceutically usable derivatives, solvates and stereoisomers thereof, including mixtures thereof in all ratios, and optionally excipients and/or adjuvants~~ and a pharmaceutically

acceptable carrier.

**Claim 20. (Currently Amended)** ~~Medicaments~~ A pharmaceutical composition comprising at least one compound of the formula-I according to Claim 1 and/or pharmaceutically usable derivatives, solvates and stereoisomers thereof, including mixtures thereof in all ratios, and at least one further medicament active ingredient an excipient, adjuvant, or vitamin.

**Claim 21. (Withdrawn-Currently Amended)** ~~Use of compounds according to Claim 1 and/or physiologically acceptable salts and solvates thereof for the preparation of a medicament~~ A method for the treatment of thromboses, myocardial infarction, arteriosclerosis, inflammation, apoplexy, angina pectoris, restenosis after angioplasty, claudicatio intermittens, migraine, tumours, tumour diseases and/or tumour metastases comprising administering to a subject in need thereof a compound of claim 1.

**Claim 22. (Currently Amended)** ~~Set (kit) consisting of separate packs of~~ A set or a kit comprising

- (a) an effective amount of a compound of the formula-I according to Claim 1 and/or pharmaceutically usable derivatives, solvates and stereoisomers thereof, including mixtures thereof in all ratios,  
and
- (b) an effective amount of a further medicament active ingredient an excipient, adjuvant, or vitamin.

**Claim 23. (Withdrawn)** ~~Use of compounds according to Claim 1 and/or physiologically acceptable salts and solvates thereof for the preparation of a medicament~~ A method for the treatment of thromboses, myocardial infarction, arteriosclerosis, inflammation, apoplexy, angina pectoris, restenosis after angioplasty, claudicatio intermittens, migraine, tumours, tumour diseases and/or tumour metastases comprising administering to a subject in need thereof a pharmaceutical composition of claim 19. , in combination with at least one further medicament active ingredient.